

Catalog # 10-1091 AG-490

CAS# 133550-30-8
(E)-2-Cyano-3-(3,4-dihydrophenyl)-*N*-(phenylmethyl)-2-propenamide
Lot # X101130

A potent inhibitor of the JAK-2 tyrosine kinase. In acute lymphoblastic leukemia (ALL) cells, which abundantly express JAK-2, AG-490 dose-dependently blocked cell growth, induced apoptosis and inhibited DNA synthesis. Blocks the growth of all pre-B ALL cells with no effect on normal B or T cells. Does not significantly inhibit other kinases such as Lck, Lyn, Btk, Syk and Src. Reduces liver injury in LPS-induced shock.³ A useful tool for exploring the role of JAK2/STAT3 pathway in physiologic processes.⁴

- 1) Wang et al. (1999), JAK3, STAT, and MAPK signaling pathways as novel molecular targets for the tyrphostin AG-490 regulation of IL-2-mediated T cell response; J. Immunol. **162** 3897
- 2) Meydan et al. (1996), Inhibition of acute lymphoblastic leukaemia by Jak-2 inhibitor; Nature 379 645
- 3) Gyurkovska and Ivanovaska (2015), *Tyrosine kinase inhibitor tyrphostin AG490 reduces liver injury in LPS-induced shock*; Eur. J. Pharmacol. **751** 118
- 4) Wu et al. (2015), ROS generated during early reperfusion contribute to intermittent hypobaric hypoxia-afforded cardioprotection against postischemia-induced Ca(+2) overload and contractile dysfunction via the JAK2/STAT3 pathway; J. Mol. Cell. Cardiol. 81 150

PHYSICAL DATA

Molecular Weight: 294.31 Molecular Formula: C₁₇H₁₄N₂O₃

Purity: 98% by TLC [10% methanol/methylene chloride, R_f = 0.50]

NMR: (Conforms)

Solubility: DMSO (up to 30 mg/mg), DMF (up to 40 mg/ml), ethanol (up to 10 mg/ml)

Physical Description: Tan solid

Storage and Stability: Store as supplied at -20°C for up to 1 year from the date of purchase. Solutions in

DMSO, DMF or ethanol may be stored at -20°C for up to 1 month.

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